



PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: J.C. Hunt, *et al.*

Serial No.: 10/086,409

Filed: November 19, 2001

For: MOUSE MONOCLONAL ANTIBODY
(5-21-3) TO HUMAN
IMMUNODEFICIENCY VIRUS GP41
PROTEIN

Attorney Docket No.: 4573.US.C6

Examiner: Not Yet Assigned

Group Art Unit: 1642

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Kimberly A. Iorio
Kimberly A. Iorio

TRANSMITTAL LETTER

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Dear Sir:

Enclosed herewith for the patent application identified above entitled MOUSE MONOCLONAL ANTIBODY (5-21-3) TO HUMAN IMMUNODEFICIENCY VIRUS GP41 PROTEIN are the following:

1. Response to Notice to Comply (1 Page), in duplicate;
2. Copy of Notice to Comply (1 Page);
3. Substitute Paper Copy of the Sequence Listing (3 Pages);
4. Substitute Computer Readable Form Copy (1 Diskette);
5. Statement to Support Filings and Submissions in Accordance with 37 CFR §§1.821-1.825 (1 Page);
6. Return Receipt Postcard.

The Commissioner is hereby authorized to charge any additional Filing Fees required under 37 CFR §1.16, as well as any patent application processing fees under 37 CFR §1.17 associated with this communication for which full payment had not been tendered, to Deposit Account No. 01-0025. A duplicate copy of this sheet is enclosed.



23492

ABBOTT LABORATORIES
Telephone: (847) 938-3137
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Respectfully submitted,
J.C. Hunt *et al.*

Dianne Casuto
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PATENT

#16

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: J.C. Hunt, *et al.*

Serial No.: 10/086,409

Filed: November 19, 2001

For: MOUSE MONOCLONAL ANTIBODY
(5-21-3) TO HUMAN
IMMUNODEFICIENCY VIRUS GP41
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Kimberly A. Iorio 9-3-02
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**RESPONSE TO NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT
APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID
SEQUENCE DISCLOSURES**

U.S. Patent and Trademark Office
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Arlington, VA 22202

Dear Sir:

In response to the Notice to Comply with Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures, dated August 22, 2002, in the patent application identified above, Applicants submit herewith a substitute Sequence Listing in computer readable form. For the Examiner's convenience, a paper copy of the Sequence Listing is also included herewith. It is also respectfully requested that the enclosed computer readable form of the Sequence Listing and corresponding paper copy replace all previously filed computer readable forms and paper copies of the Sequence Listing.

I hereby state that the content of the computer readable copy of the Sequence Listing submitted herewith is identical to the paper copy of the Sequence Listing submitted herewith, and that the computer readable copy contains no sequence information that would constitute new matter beyond the originally filed application.

The Commissioner is hereby authorized to charge the payment of any patent application processing fees under 37 CFR 1.17 concerning this transaction, or to credit any overpayment to Deposit Account No. 01-0025. A duplicate copy of this sheet is attached.



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Dianne Casuto

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Attorney for Applicants



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APPLICATION NUMBER	FILING/RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
10/086,409	11/19/2001	Jeffrey C. Hunt	4573.US.C6

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CONFIRMATION NO. 7509

FORMALITIES LETTER



OC000000008674094

Date Mailed: 08/22/2002

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant is given **TWO MONTHS FROM THE DATE OF THIS NOTICE** within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

- The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d). Applicant must provide a substitute computer readable form (CRF) copy of the "Sequence Listing" and a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d). If applicant desires the sequence listing in the instant application to be identical with that of another application on file in the U.S. Patent and Trademark Office, such request in accordance with 37 CFR 1.821(e) may be submitted in lieu of a new CRF.

For questions regarding compliance to these requirements, please contact:

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PART 2 - COPY TO BE RETURNED WITH RESPONSE